UNITED NATIONS

E/NL.1987/73-74

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LAWS AND REGULATIONS

PROMULGATED TO GIVE EFFECT TO THE PROVISIONS OF THE INTERNATIONAL TREATIES ON NARCOTIC DRUGS AND PSYCHOTROPIC SUBSTANCES

In accordance with the relevant articles of the international treaties on narcotic drugs and psychotropic substances, the Secretary-General has the honour to communicate the following legislative texts.

FEDERAL REPUBLIC OF GERMANY

Communicated by the Government of the Federal Republic of Germany

NOTE BY THE SECRETARIAT

(a) Some editing of texts may be done by the Secretariat in the interest of clarity. In this connection, words in square brackets [ ] have been added or changed by the Secretariat.

(b) Only passages directly relevant to the control of narcotic drugs or psychotropic substances have been reproduced in this document. Non-relevant parts of laws and regulations have been deleted by the Secretariat; such deletions are indicated by [...].

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* Note by the Secretariat: The present document is a direct reproduction of the texts communicated to the Secretariat.

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E/NL.1987/73

First Ordinance
Amending the Regulations of Narcotic Drugs Legislation
(First Narcotic Drugs Legislation Amending Ordinance)
6 August 1984

On the basis of section 1 paragraph 2 and section 13 paragraph 3 of the Narcotics Act of 28 July 1981 (Federal Law Gazette I, p. 681, 1187) the Federal Government, with the consent of the Bundesrat, hereby orders as follows:

Article 1

Amendment of the Schedules of the Narcotics Act

The Schedules of the Narcotics Act shall be amended as follows:

1. Schedule I of the Narcotics Act shall be amended as follows:

   a) The narcotic drugs etorphine and phendimetrazine shall be deleted with all particulars given.

   b) The following narcotic drugs shall be inserted in alphabetical order:

"Dimethoxybromampheta=mine (DOB)	2,5-Dimethoxy-4-bromamphetamine
Ethylpiperidyl=benzilate (JB 318)	1-Ethyl-3-piperidylbenzilate
Methoxyamphetamine (PMA)	4-Methoxy-α-methylphenethylamine
Methoxymethylene-dioxyamphetamine (MMDA)	3-Methoxy-α-methyl-4,5-methylene-dioxyphenethylamine
Methylenedioxyamphet-amine (MDA)	α-Methyl-3,4-methylenedioxyphene=thylamine
Methylpiperidyl=benzilate (JB 336)	1-Methyl-3-piperidylbenzilate
Racemethorphan	(±)-3-Methoxy-17-methylmorphinan
Trimethoxyamphetamine (TMA)	3,4,5-Trimethoxy-α-methyl-phenethylamine".

   c) The scientific designation of the narcotic drug etoxeridine shall be indicated in the following version:

   "Ethyl-{1-[2-(2-hydroxyethoxy)ethyl]-4-phenyl-4-piperidin-carboxylate}".

2. Schedule II of the Narcotics Act shall be amended as follows:

a) The narcotic drug papaver somniferum shall be deleted with all particulars given.

b) The following narcotic drugs shall be inserted in alphabetical order:

"Cetobemidon	1-/4-(3-Hydroxyphenyl)-1-methyl-4-piperidyl/-1-propanon
Dextropropoxyphene	(+)-(1-Benzyl-3-dimethylamino-2-methyl-1-phenylpropyl)propionate - except in preparations containing without any additional substance listed in Schedules I to III, up to 2.5 per cent in oral application or, per dosage unit, up to 135 milligrammes of dextropropoxyphene, calculated as a base -
Levorphanol	(-)-17-Methyl-3-morphinanol
Phendimetrazine	3,4-Dimethyl-2-phenylmorpholine".

c) The position of the narcotic drug papaver orientale (papaver bracteatum) shall be indicated in the following version:

"Papaver orientale (Papaver bracteatum)	plants and plant segments of plants belonging to the species of papaver orientale, except their seeds; if they serve ornamental purposes, narcotic drug regulations shall only apply to import, export and transit".

3. Schedule III Part A of the Narcotics Act shall be amended as follows:

a) The following narcotic drugs shall be deleted with all particulars given:

Cetobemidon, dextropropoxyphene, levorphanol.

b) The position of the narcotic drug tilidine shall be indicated in the following version:

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"Tilidine	Ethyl-(2-dimethylamino-1-phenyl-3-cyclohexen-1-carboxylate) - except in preparations containing without any additional substance listed in Schedules I to III, up to 615 milligrammes of tilidine per dosage unit, calculated as a base and related to this quantity, not less than 7.5 per cent of naloxone-hydrochloride. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -".

c) The following narcotic drugs shall be inserted in alphabetical order:

"Alfentanil	N-{1-[2-(4-ethyl-5-oxo-2-tetrazolin-1-yl)ethyl-4-methoxymethyl-4-piperidyl} propionanilide
Buprenorphin	17-Cyclopropylmethyl-4,5α-epoxy-7α-[(S)-1-hydroxy-1,2,2-trimethylpropyl]-6-methoxy-6,14-endo-ethanomorphinan-3-ol
Etorphine	4,5α-Epoxy-7α-(1-hydroxy-1-methylbutyl)-6-methoxy-17-methyl-6,14-endo-ethenomorphinan-3-ol
Nabilone	(±)-trans-3-(1,1-Dimethylheptyl)-7,8,10,10a-tetrahydro-1-hydroxy-6,6-dimethyl-6H-dibenzo[b,d]pyran-9(6aH)-on
Papaver somniferum	plants and plant segments of plants belonging to the species of papaver somniferum, except their seeds; if they serve ornamental purposes and if morphine has been extracted by a method authorized by the Federal Health Office, narcotic drug regulations shall only apply to import, export, transit, cultivation and production".

4. Schedule III Part B shall be amended as follows:

a) The following narcotic drug shall be inserted in alphabetical order:

"Pentazocine	1,2,3,4,5,6-hexahydro-6,11-dimethyl-3(3-methyl-2-butenyl)-2,6-methano-3-benzazocin-8-ol".

b) The scientific designation of the narcotic drug pentobarbital shall be indicated in the following version:

"5-Ethyl-5-(1-methylbutyl)-barbituric acid".

5. In Schedule III Part C the exceptions for the narcotic drug barbital shall be amended by the following wording:

"- except in preparations which without being administered to human or animal bodies exclusively serve diagnostic or analytic purposes and which per dosage unit contain up to 25 g barbital or barbital sodium or a mixture of these two substances -".

Article 2

Trade in New Narcotic Drugs and Exempted Preparations

(1) A person who on coming into force of this ordinance without belonging to the group of persons mentioned in section 4 of the Narcotics Act participates in the trade in narcotic drugs (section 3 paragraph 1 (1) of the Narcotics Act) or in the trade in exempted preparations (section 3 paragraph 1 (2) of the Narcotics Act) with the substances mentioned below:

1. Alfentanil
2. Buprenorphin
3. Dimethoxybromoamphetamine (DOB)
4. Ethylpiperidylø-benzilate (JB 318)
5. Methoxyamphetamine (PMA)
6. Methoxymethylenedioxyamphetamine (MMDA)
7. Methylenedioxyamphetamine (MDA)
8. Methylpiperidylø-benzilate (JB 336)
9. Nabilon
10. Pentazocine
11. Racemethorphan
12. Trimethoxyamphetamine (TMA)

shall be entitled to do so up to 31 December 1984. In case he applies for a licence pursuant to section 3 paragraph 1 of the Narcotics Act before 1 January 1985 this entitlement shall continue until the application is dismissed finally and absolutely.

(2) A person who at the time this ordinance comes into force, as a person entitled within the meaning of the 1st sentence of paragraph 1, has in safekeeping narcotic drugs as described in the a.m. paragraph shall be under the obligation by 31 December 1984

1. to report to the Federal Health Office these narcotic drugs indicating kind and amount and
2. to hand them over or sale them to the holder of a licence pursuant to section 3 paragraph 1 of the Narcotics Act, to the operator of a pharmacy or a veterinary house dispensary or to a person untitled pursuant to the 2nd sentence of paragraph 1 or to destroy them under the terms of section 16 of the Narcotics Act if he does not file an application pursuant to the 2nd sentence of paragraph 1.

A person who according to sentence 1 (2) acquires narcotic drugs shall have to inform the Federal Health Office by 31 March 1985 on the dispensing party as well as kind and quantity of the narcotic drug acquired.

(3) If the narcotic drugs specified in paragraph 1 are packed for dispensing to the consumer without the packages complying with the requirements pursuant to section 14 of the Narcotics Act, they shall still be allowed to be dispensed in these packages until 31 July 1985.

(4) If the narcotic drugs specified in paragraph 1 are not stored and secured in the manner required in section 15 of the Narcotics Act, they shall still be allowed to be stored in the manner admissible so far until 31 July 1985. The 1st sentence shall not apply to storage in pharmacies, veterinary house dispensaries and on merchant ships.

Article 3

Amendment of the Narcotic Drugs Prescription Order

The Narcotic Drugs Prescription Order of 16 December 1981 (Federal Law Gazette I, p. 1427) shall be amended as follows:

1. Section 2 paragraphs 1 to 3 shall be worded as follows:

"(1) A medical practitioner shall be allowed to prescribe for a patient on any single day:

a) one of the following narcotics in compliance with maximum quantities established below and other restrictions concerning purpose of application, content and dosage form (other restrictions):

1.	Amphetamine	200 mg
2.	Buprenorphin	4 mg
3.	Dextromoramide	100 mg
4.	Hydrocodone	200 mg
5.	Hydromorphone	30 mg
6.	Levomethadone	60 mg
7.	Methamphetamine	100 mg
8.	Methaqualone	6,000 mg
9.	Methylphenidate	200 mg
10.	Morphine	200 mg
11.	Nabilon	36 mg
12.	Normethadone	200 mg
13.	Opium, standardized	2,000 mg
14.	Opium extract	1,000 mg
15.	Tincture of opium	20,000 mg
16.	Native opium alkaloids in their natural mixing ratio, calculated as morphine	200 mg
17.	Oxycodone	200 mg
18.	Papaver somniferum, calculated as morphine	200 mg
19.	Pentazocine	700 mg
20.	Pethidine	1,000 mg
21.	Phenmetrazine	600 mg
22.	Piritramide	220 mg
23.	Thebacon	200 mg
24.	Tilidine	1,050 mg

or

b) a preparation containing one or two narcotics specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C of the Narcotics Act.

(2) If in cases of particularly severe illnesses the quantity to be administered to the patient requires the maximum quantity established under paragraph 1 (a) to be exceeded, the practitioner shall be allowed to prescribe for this patient on any single day:

1. one of the narcotic drugs specified in paragraph 1 (a) nos 2,3,5,6,10,13 to 20,22 and 24, up to four times the established maximum quantity or

2. if the patient is under long-term treatment, the narcotic drug specified in paragraph 1 (a) no 10, up to ten times the established maximum quantity.

In such cases he shall have to provide the prescription with a hand-written note saying "quantity medically justified". A prescription according to point 2 shall only be admissible for oral application as a solution up to a content of 4 per cent adding at least 0.05 per cent quinine hydrochloride and 1 per cent carboxymethylcellulose-sodium; it shall have to be provided with directions for use indicating that the solution is intended "only for oral application" and at what intervals it has to be taken. Notwithstanding the first sentence he shall be allowed to prescribe the solution specified in the third sentence in the quantity required for three days.

(3) As his own surgery supply a medical practitioner shall be allowed only to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 up to the established maximum quantity or a preparation containing one or two of the narcotic drugs specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C of the Narcotics Act,

2. up to 30 mg alfentanil for pre-medication and anaesthesia including neuroleptoanalgesia,

3. up to 1,000 mg cocaine for surgeries on the eye, larynx, nose, ear, throat or jaw as

   a) solution up to a content of 20 per cent, or

   b) eye tablet or ointment up to a content of 2 per cent,

4. up to 10 mg fentanyl for pre-medication and anaesthesia including neuroleptoanalgesia, for diagnostic surgeries and in intensive care medicine and

5. up to 2,500 mg pentobarbital for pre-medication and anaesthesia".

2. In section 2 paragraph 4 the quotation "nos 2 and 3" shall be replaced by the quotation "nos 2 to 5".

3. Section 3 paragraphs 1 and 2 shall be worded as follows:

"(1) A dental surgeon shall be allowed to prescribe for a patient on any single day:

a) one of the following narcotic drugs in compliance with the maximum quantities established below:

1.	Amphetamine	200 mg
2.	Buprenorphin	4 mg
3.	Dextromoramide	100 mg
4.	Hydrocodone	200 mg
5.	Hydromorphone	30 mg
6.	Levomethadone	60 mg
7.	Methamphetamine	100 mg
8.	Methaqualone	6,000 mg
9.	Methylphenidate	200 mg
10.	Morphine	200 mg
11.	Normethadone	200 mg
12.	Opium, standardized	2,000 mg
13.	Opium extract	1,000 mg
14.	Opium tincture	20,000 mg
15.	Native opium alkaloids in their natural mixing ratio, calculated as morphine	200 mg
16.	Oxycodone	200 mg

17. Pentazocine	700 mg
18. Pethidine	1,000 mg
19. Phenmetrazine	600 mg
20. Piritramide	220 mg
21. Thebacon	200 mg
22. Tilidine	1,050 mg

or

b) a preparation containing one or two of the narcotic drugs specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C of the Narcotics Act.

(2) As his own surgery supply a dental surgeon shall only be allowed to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 up to the established maximum quantity or a preparation containing one or two of the narcotic drugs specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C of the Narcotics Act,
2. up to 30 mg alfentanil for pre-medication and anaesthesia including neuroleptoanalgesia,
3. up to 5 mg fentanyl for pre-medication and anaesthesia including neuroleptoanalgesia and
4. up to 2,500 mg pentobarbital for pre-medication and anaesthesia".

4. In section 3 paragraph 3 the quotation "no 2" shall be replaced by the quotation "nos 2 to 4".

5. Section 4 paragraphs 1 to 3 shall be worded as follows:

"(1) A veterinary surgeon shall be allowed to prescribe for an animal on any single day:

a) one of the following narcotic drugs in compliance with the maximum quantities established below:

1. Amphetamine	1,000 mg
2. Buprenorphin	4 mg

3.	Dextromoramide	100 mg
4.	Hydrocodone	200 mg
5.	Hydromorphone	30 mg
6.	Levomethadone	250 mg
7.	Methamphetamine	100 mg
8.	Methaqualone	6,000 mg
9.	Methylphenidate	200 mg
10.	Morphine	500 mg
11.	Normethadone	200 mg
12.	Opium, standardized	12,000 mg
13.	Opium extract	6,000 mg
14.	Opium tincture	120,000 mg
15.	Native opium alkaloids in their natural mixing ratio, calculated as morphine	1,500 mg
16.	Oxycodone	300 mg
17.	Pentazocine	700 mg
18.	Pethidine	1,000 mg
19.	Phenmetrazine	600 mg
20.	Piritramide	220 mg
21.	Thebacon	200 mg
22.	Tilidine	1,050 mg

or

b) a preparation containing one or two of the narcotic drugs specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C of the Narcotics Act.

(2) If in cases of particularly severe illnesses the quantity to be administered to the animal requires the maximum quantity established in paragraph 1 (a) to be exceeded, the veterinary surgeon shall be allowed to prescribe for this animal on any single day only one of the narcotic drugs specified in paragraph 1 (a) nos 2,3,5,6,10,12 to 18, 20 and 22 up to twice the established quantity. In such cases he shall have to provide the prescription with a hand-written note saying "quantity veterinarily justified".

(3) As his own surgery supply a veterinary surgeon shall only be allowed to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 up to the established maximum quantity or a preparation containing one or two of the narcotic drugs specified in Schedule III Part B, except pentazocine and pentobarbital, or Part C' of the Narcotics Act,

2. up to 30 mg alfentanil for pre-medication and anaesthesia including neuroleptoanalgesia,

3. up to 1,000 mg cocaine for surgeries on the eye as

   a) solution up to a content of 20 per cent, or

   b) eye tablet or ointment up to a content of 2 per cent,

4. up to 30 mg etorphine for immobilization of zoo and wild animals,

5. up to 10 mg fentanyl for pre-medication and anaesthesia including neuroleptoanalgesia, for diagnostic surgeries and for immobilization and

6. up to 40,000 mg pentobarbital for pre-medication and anaesthesia as well as for putting down animals".

6. In section 4 paragraph 4 the quotation "nos 2 and 3" shall be replaced by the quotation "nos 2 to 6".

7. In section 6 paragraph 1 the sub-paragraphs 4, 5 and 7 shall be worded as follows:

"4. directions for use specifying the single and the daily dose; in the cases contained in section 2 paragraph 2 (2) also the intervals of application,

5. for prescriptions of alfentanil, cocaine, etorphine, fentanyl and pentobarbital the purpose of administration,

7. in the cases referred to in section 2 paragraph 2 and section 4 paragraph 2 the annotations stipulated thereunder."

8. Section 8 paragraph 2 sentence 2 shall be worded as follows:

"He shall be allowed to prescribe for merchant ships

1. having no ship's doctor only the narcotic drugs specified in section 2 paragraph 1 (a) nos 5, 13 and 20,

2. having a ship's doctor only the narcotic drugs specified in section 2 paragraph 1 (a) nos 4,5,7,10,13,17 and 20."

9. Section 10 sentence 1 (2)(a) shall be worded as follows:

"a) contrary to section 2 paragraphs 1, 2 sentences 1, 3 or 4 or section 3 paragraph 1 for a patient,".

Article 4

Berlin Clause

Pursuant to section 14 of the Third Transitory Law in conjunction with section 41 sentence 2 of the Narcotics Act this order shall also apply in the Land of Berlin.

Article 5

Entry into Force, Transitory Provision, Replaced Provisions

(1) This Ordinance shall enter into force on 1 September 1984.

(2) Plants or plant segments of plants belonging to the species of papaver somniferum, if they serve ornamental purposes and if morphine has not been extracted, shall until 31 March 1985 be allowed to be marketed under the previous provisions by persons who have neither imported nor cultivated them.

(3) The Ninth Ordinance Concerning Substances Placed on the Same Footing as Narcotic Drugs of 15 June 1981 (Federal Law Gazette I, p. 529) shall be repealed.

Bonn, 6 August 1984

For the Federal Chancellor
The Federal Minister of Posts and Telecommunications
Dr. Christian Schwarz-Schilling

The Federal Minister for Youth,
Family Affairs and Health
Geißler

E/NL.1987/74

Second Ordinance
Amending the Regulations of Narcotic Drugs Legislation
(Second Narcotic Drugs Legislation Amending Ordinance)
23 July 1986

On the basis of section 1 paragraph 2 of the Narcotics Act of 28 July 1981 (Federal Law Gazette I, p. 681, 1187) the Federal Government, having heard experts, hereby orders as follows,

on the basis of section 1 paragraph 3 of the Narcotics Act the Federal Minister for Youth, Family Affairs, Women and Health hereby orders as follows,

on the basis of section 13 paragraph 3 of the Narcotics Act the Federal Government, with the consent of the Bundesrat, hereby orders as follows:

Article 1

Amendment of the Narcotics Act

The Narcotics Act amended by article 1 of the ordinance of 6 August 1984 (Federal Law Gazette, p.1081) shall be amended as follows:

1. In Schedule I the following narcotic drugs shall be inserted in alphabetical order:

Cathinone	(S)-2-Aminopropiophenone
Dimethoxyamphetamine (DMA)	2,5-Dimethoxy-α-methylphenethylamine
Dimethoxyethylamphetamine (DOET)	4-Ethyl-2,5-dimethoxy-α-methylphenethylamine
Methylendioxymethamphetamine (MDMA)	N,α-Dimethyl-3,4-methylendioxyphenethylamine

2. Schedule II shall be amended as follows:
   a) The position of the narcotic drug papaver orientale (papaver bracteatum) shall be indicated in the following version:

"Papaver bracteatum — plants and plant segments of plants belonging to the species of papaver bracteatum, except their seeds, - except for ornamental purposes -"

b) The following narcotic drug shall be inserted in alphabetical order:

Thebacon — 4,5 α-Epoxy-3-methoxy-17-methyl-6-morphinen-6-ylacetate

3. Schedule III, Part A shall be amended as follows:

a) The position of the narcotic drug opium shall be indicated in the following version:

"Opium — the coagulated juice of plants belonging to the species of papaver somniferum

- except in preparations manufactured according to a homoeopathic processing method in particular according to the homoeopathic part of the pharmacopoeia if the final concentration does not exceed the sixth degree of dilution -."

b) The position of the narcotic drug papaver somniferum shall be indicated in the following version:

"Papaver somniferum — plants and plant segments of plants belonging to the species of papaver somniferum, except their seeds

- except for plants and plant segments cultivated for ornamental purposes (poppy straw) if morphine has been extracted by a method authorized by the Federal Health Office; in this case the narcotic drug regulations shall only apply to import, export and transit -

- except in preparations manufactured according to a homoeopathic processing method in particular according to the homoeopathic part of the pharmacopoeia if the final concentration does not exceed the fourth degree of dilution -."

c) The narcotic drug thebacon shall be deleted with all particulars given.

d) The following narcotic drug shall be inserted in alphabetical order:

Fenetylline	7-[2-[(α-Methylphenetyl)amino]ethyl]theophilline

4. In Schedule III Part C the following narcotic drugs shall be inserted in alphabetical order:

Alprazolam	8-Chloro-1-methyl-6-phenyl-4H-[5]triazolo[4,3-a][1,4]benzodiazepine

- except in preparations containing without any additional substance listed in Schedules I to III, up to 1 mg alprazolam per dosage unit. These preparations, howeve are subject to the narcotic drugs regulations governing import, export and transit -

Bromazepam	7-Bromo-1,3-dihydro-5-(2-pyridyl)-2H-1,4-benzodiazepin-2-one

- except in preparations containing without any additional substance listed in Schedules I to III, up to 12 mg bromazepam per dosage unit. These preparations,however are subject to the narcotic drugs regulations governing import, export and transit -

Camazepam	(7-Chloro-1,3-dihydro-3-hydroxy-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one) dimethylcarbamate(ester) - except in preparations containing without any additional substance listed in Schedules I to III, up to 20 mg camazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Chlordiazepoxide	7-Chloro-2-methylamino-5-phenyl-3H-1,4-benzodiazepine-4-oxide - except in preparations containing without any additional substance listed in Schedules I to III, up to 40 mg chlordiazepoxide per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Chlobazam	7-Chloro-1-methyl-5-phenyl-1H-1,5-benzodiazepine-2,4(3H,5H)-dione - except in preparations containing without any additional substance listed in Schedules I to III, up to 30 mg chlobazam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Clonazepam	5-(o-Chlorophenyl)-1,3-dihydro-7-nitro-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 0.25 per cent as drip solution, but not more than 250 mg per package unit or up to 2 mg clonazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Clorazepate	7-Chloro-2,3-dihydro.2,2-dihydroxy-5-phenyl-1H-1,4-benzodiazepine-3-carboxylic acid - except in preparations containing without any additional substance listed in Schedules I to III, up to 50 mg per dosage unit, as a dry substance for parenteral application only up to 100 mg;dipotassium-clorazepate. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -

Clotiazepam	5-(o-Chlorophenyl)-7-ethyl-1,3-dihydro-methyl-2 H-thieno [2,3-e] [1,4] diazepin-2-one -except in preparations containing without any additional substance listed in Schedules I to III, up to 20 mg clotiazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Cloxazolam	10-Chloro-11b-(o-chlorophenyl)-2,3,7,11b-tetrahydro-oxazolo [3,2-d] [1,4] benzodiazepin-6(5H)-one
Delorazepam	7-Chloro-5-(o-chlorophenyl)-1,3-dihydro-2H-1,4-benzodiazepin-2-one
Diazepam	7-Chloro-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one -except in preparations containing without any additional substance listed in Schedules I to III, up to 1 per cent as syrup or drip solution but not more than 250 mg per package unit or up to 20 mg diazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Estazolam	8-Chloro-6-phenyl-4H-5-triazolo [4,3-a] [1,4] benzodiazepine
Ethylloflazepate	Ethyl-7-chloro-5-(o-fluorophenyl)-2,3-dihydro-2-oxo-1H-1,4-benzodiazepin-3-carboxylate
Fludiazepam	7-Chloro-5-(o-fluorophenyl)-2,3-dihydro-1-methyl-1H-1,4-benzodiazepin-2-one
Flunitrazepam	5-(o-Fluorophenyl)-2,3-dihydro-1-methyl-7-nitro-1H-1,4-benzodiazepin-2-one -except in preparations containing without any additional substance listed in Schedules I to III, up to 2 mg flunitrazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Flurazepam	7-Chloro-1-[2-(diethylamino)ethyl]-5-(o-fluorophenyl)-1,3-dihydro-2H-1,4-benzodiazepin-2-one -except in preparations containing without any additional substance listed in

	Schedules I to III, up to 30 mg flurazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Halazepam	7-Chloro-1,3-dihydro-5-phenyl-1-(2,2,2-trifluoroethyl)-2H-1,4-benzodiazepin-2-one -except in preparations containing without any additional substance listed in Schedules I to III, up to 120 mg halazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Haloxazolam	10-Bromo-11b-(o-fluorophenyl)-2,3,7,11b-tetrahydrooxazolo[3,2-d][1,4] benzodiazepin-6(5H)-one
Ketazolam	11-Chloro-8,12b-dihydro-2,8-dimethyl-12b-phenyl-4H-[1,3]oxazino[3,2-d][1,4] benzodiazepin-4,7(6H)-dione -except in preparations containing without any additional substance listed in Schedules I to III, up to 45 mg ketazolam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Loprazolam	6-(o-Chlorophenyl)-2,4-dihydro (4-methyl-1-piperazinyl)methylene -8-nitro-1H-imidazo[1,2-a][1,4] benzodiazepin-1-one - except in preparations containing without any additional substance listed in Schedules I to III up to 2.5 loprazolam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Lorazepam	7-Chloro-5-(o-chlorophenyl)-1,3-dihydro-3-hydroxy-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 2.5 mg lorazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Lormetazepam	7-Chloro-5-(o-chlorophenyl)-1,3-dihydro-3-hydroxy-1-methyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without

	any additional substance listed in Schedules I to III, up to 2 mg lormetazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Medazepam	7-Chloro-2,3-dihydro-1-methyl-5-phenyl-1H-1,4-benzodiazepine - except in preparations containing without any additional substance listed in Schedules I to III, up to 10 mg medazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Nimetazepam	1,3-Dihydro-1-methyl-7-nitro-5-phenyl-2H-1,4-benzodiazepin-2-one
Nitrazepam	1,3-Dihydro-7-nitro-5-phenyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III up to 0.5 per cent as drip solution but not more than 250 mg per package unit or up to 10 mg nitrazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Nordazepam	7-Chloro-1,3-dihydro-5-phenyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III up to 0.5 per cent as drip solution but not more than 150 mg per package unit or up to 15 mg nordazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Oxazepam	7-Chloro-1,3-dihydro-3-hydroxy-5-phenyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 50 mg oxazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -

Oxazolam	10-Chloro-2,3,7,11b-tetrahydro-2-methyl-11b-phenyloxazolo [3,2-d] [1,4]benzodiazepin-6(5H)-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 20 mg oxazolam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Pinazepam	7-Chloro-1,3-dihydro-5-phenyl-1-(2-propinyl)-2H-1,4-benzodiazepin-2-one
Prazepam	7-Chloro-1-(cyclopropylmethyl)-1,3-dihydro--5-phenyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 20 mg prazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Temazepam	7-Chloro-1,3-dihydro-3-hydroxy-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 20 mg temazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Tetrazepam	7-Chloro-5-(1-cyclohexen-1-yl)-1,3-dihydro-1-methyl-2H-1,4-benzodiazepin-2-one - except in preparations containing without any additional substance listed in Schedules I to III, up to 50 mg tetrazepam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -
Triazolam	8-Chloro-6-(o-chlorophenyl)-1-methyl-4H-5-triazolo[4,3-a] [1,4]benzodiazepine - except in preparations containing without any additional substance listed in Schedules I to III, up to 1 mg triazolam per dosage unit. These preparations, however, are subject to the narcotic drugs regulations governing import, export and transit -

5. After section 40 the following section 40 (a) shall be inserted:

"Section 40 a

Trade in Further New Narcotic Drugs
and Exempted Preparations

(1) A person who on 31 July 1986 without belonging to the group of persons mentioned in section 4 of the Narcotics Act participates in the trade in narcotic drugs (section 3 paragraph 1 (1) or in the trade in exempted preparations (section 3 paragraph 1 (2) with the below mentioned substances their isomers, esters, ethers, molecular compounds and salts:

1. Alprazolam
2. Bromazepam
3. Camazepam
4. Chlordiazepoxide
5. Clobazam
6. Clonazepam
7. Clorazepate
8. Clotiazepam
9. Cloxazolam
10. Delorazepam
11. Diazepam
12. Estazolam
13. Ethylloflazepate
14. Fenetylline
15. Fludiazepam
16. Flunitrazepam
17. Flurazepam
18. Halazepam
19. Haloxazolam
20. Ketazolam
21. Loprazolam
22. Lorazepam
23. Lormetazepam
24. Medazepam
25. Nimetazepam
26. Nitrazepam
27. Nordazepam
28. Oxazepam
29. Oxazolam
30. Pinazepam
31. Prazepam
32. Temazepam
33. Tetrazepam
34. Triazolam

shall be entitled to do so up to 31 October 1986. In case he applies for a licence pursuant to section 3 paragraph 1 before 1 November 1986 this entitlement shall continue until the application is dismissed finally and absolutely. The person entitled pursuant to sentence 1 or 2 shall, like the holder of a licence, as from the time of this ordinance coming into force, be bound by all provisions of this act and the ordinances issued in connection therewith.

(2) If the narcotic drugs specified in paragraph 1 are packed for dispensing to the consumer without the packages complying with the requirements pursuant to section 14, they shall still be allowed to be dispensed in these packages until 30 June 1987.

(3) If the narcotic drugs specified in paragraph 1 are not stored and secured in the manner required in section 15, they shall still be allowed to be stored in the manner admissible so far until 30 June 1987. The ist sentence shall not apply to storage in pharmacies, veterinary house dispensaries and on merchant ships."

Article 2

Amendment of the Narcotic Drugs Prescription Ordinance

The Narcotic Drugs Prescription Ordinance of 16 December 1981 (Federal Law Gazette I, p. 1427) amanded by article 3 of the ordinance of 6 August 1984 (Federal Law Gazette I, p. 1081) shall be amended as follows:

1. Section 2 shall be worded as follows:

"Section 2

Prescribing by a medical practitioner

(1) A medical practitioner shall be allowed to prescribe for a patient on any single day:

a) one of the following narcotics in compliance with the maximum quantities established below (single maximum quantities):

1.	Amphetamine	200 mg
2.	Buprenorphin	10 mg
3.	Dextromoramide	100 mg
4.	Fenetylline	2,500 mg
5.	Hydrocodone	200 mg
6.	Hydromorphone	30 mg
7.	Levomethadone	60 mg
8.	Methamphetamine	100 mg
9.	Methaqualone	6,000 mg
10.	Methylphenidate	200 mg
11.	Morphine	200 mg
12.	Nabilone	36 mg
13.	Normethadone	200 mg
14.	Opium, standardized	2,000 mg
15.	Opium extract	1,000 mg

16. Tincture of opium	20,000 mg
17. Oxycodone	200 mg
18. Papaver somniferum, calculated as morphine	200 mg
19. Pentazocine	700 mg
20. Pentobarbital	2,500 mg
21. Pethidine	1,000 mg
22. Phenmetrazine	600 mg
23. Piritramide	220 mg
24. Tilidine	1,050 mg

or

b) one of the narcotic drugs specified in Schedule III Part B with the exception of pentazocine and pentobarbital or Part C of the Narcotics Act.

(2) In a case of a particularly severe illness, if the severeness of the illness requires it, a medical practitioner shall be allowed to prescribe on any single day for a patient whom he treats on a long-term basis, one of the below-mentioned narcotic drugs in the following manner:

1. Buprenorphin, dextromoramide, hydromorphone, levomethadone, morphine, standardized opium, opium extract, tincture of opium, oxycodone, pentazocine or pethidine up to twice the single maximum quantity in the amount required for up to 7 days,

2. Morphine for oral application only
   a) as capsules or tablets with retarded active substance release per day of application up to five times,
   b) as a solution up to a content of 4 per cent with at least 1 per cent carboxymethylcellulose-sodium being added, per day of application up to ten times

   the single maximum quantity in the amount required for up to 7 days or

3. Buprenorphin or morphine for peridural and intrathecal application per day of application up to the single maximum quantity in the amount required for up to 28 days.

Narcotic drug dependence shall not be considered as a case of a particularly severe illness within the meaning of the first sentence.

(3) As his own surgery supply the medical practitioner shall be allowed only to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 for up to the established maximum quantity or one of the narcotic drugs specified in paragraph 1 (b),

2. up to 275 mg alfentanil for pre-medication and anaesthesia including neurolepto- and on-demand analgesia,

3. up to 1,000 mg cocaine for surgeries on the eye, larynx, nose, ear, throat or jaw as
   a) solution up to a content of 20 per cent or
   b) eye tablet or ointment up to a content of 2 per cent and

4. up to 40 mg fentanyl for pre-medication and anaesthesia including neurolepto- and on-demand analgesia, for diagnostic surgeries and intensive-care medicine.

(4) The medical practitioner who heads or supervises a partial unit (ward) of a departmentized hospital or who heads or supervises an undepartmentized hospital shall be allowed to prescribe as ward supply for the facility headed or supervised by him, only the narcotic drugs specified in the paragraphs 1 and 3 nos 2 to 4 in consideration of the restrictions pursuant to paragraph 3 as regards intended purpose, content and dosage form. This shall also apply to an out-of-hospital medical practitioner who has a certain number of hospital beds reserved for him if the beds allocated to him are separated from other partial units as regards location or organisation."

2. Section 3 shall be worded as follows:

"Section 3

Prescribing by a dental surgeon

(1) A dental surgeon shall be allowed to prescribe for a patient on any single day:

a) one of the following narcotic drugs in compliance with the maximum quantities established below (single maximum quantities):

1.	Amphetamine	200 mg
2.	Buprenorphin	10 mg
3.	Dextromoramide	100 mg
4.	Fenetylline	2,500 mg
5.	Hydrocodone	200 mg
6.	Hydromorphone	30 mg
7.	Levomethadone	60 mg
8.	Methamphetamine	100 mg
9.	Methaqualone	6,000 mg
10.	Methylphenidate	200 mg
11.	Morphine	200 mg
12.	Normethadone	200 mg
13.	Opium, standardized	2,000 mg
14.	Opium extract	1,000 mg
15.	Tincture of opium	20,000 mg
16.	Oxycodone	200 mg
17.	Pentazocine	700 mg
18.	Pethidine	1,000 mg
19.	Phenmetrazine	600 mg
20.	Piritramide	220 mg
21.	Tilidine	1,050 mg

or

b) one of the narcotic drugs specified in Schedule III Part B with the exception of pentazocine and pentobarbital or Part C of the Narcotics Act.

(2) As his own surgery supply the dental surgeon shall be allowed only to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 (a) up to the established maximum quantity or one of the narcotic drugs specified in paragraph 1 (b),
2. up to 30 mg alfentanil for pre-medication and anaesthesia including neuroleptoanalgesia,
3. up to 5 mg fentanyl for pre-medication and anaesthesia including neuroleptoanalgesia and
4. up to 2,500 mg pentobarbital for pre-medication and anaesthesia.

(3) The dental surgeon who heads or supervises a partial unit (ward) of a departmentized or who heads or supervises an undepartmentized hospital shall be allowed to prescribe as ward supply for the facility headed or supervised by him only the narcotic drugs specified in the paragraphs 1 and 2 nos 2 to 4 in consideration of the restrictions pursuant to paragraph 2 as regards the intended purpose. This shall also apply to an out-of-hospital dental surgeon who has a certain number of hospital beds reserved for him if the beds allocated to him are separated from other partial units as regards location or organisation."

3. Section 4 shall be worded as follows:

"Section 4

Prescribing by a veterinary surgeon

(1) A veterinary surgeon shall be allowed to prescribe for an animal on any single day:

a) one of the following narcotic drugs in compliance with the maximum quantities established below (single maximum quantities):

1.	Amphetamine	1,000 mg
2.	Buprenorphin	10 mg
3.	Dextromoramide	100 mg
4.	Fenetylline	2,500 mg
5.	Hydrocodone	200 mg
6.	Hydromorphone	30 mg
7.	Levomethadone	250 mg
8.	Methamphetamine	100 mg
9.	Methaqualone	6,000 mg
10.	Methylphenidate	200 mg
11.	Morphine	500 mg
12.	Normethadone	200 mg
13.	Opium, standardized	12,000 mg
14.	Opium extract	6,000 mg
15.	Opium tincture	120,000 mg
16.	Oxycodone	300 mg
17.	Pentazocine	700 mg
18.	Pethidine	1,000 mg
19.	Phenmetrazine	600 mg
20.	Piritramide	220 mg
21.	Tilidine	1,050 mg

or

b) one of the narcotic drugs specified in Schedule III Part B with the exception of pentazocine and pentobarbial or Part C of the Narcotics Act.

(2) In case of a particularly severe illness, if the severeness of the illness requires it, a veterinary surgeon shall be allowed to prescribe for an animal on any single day one of the following narcotic drugs:

Buprenorphin, dextromoramide, hydromorphone, levomethadone, morphine, opium,standardized,opium extract, opium tincture, oxycodone, pentazocine or pethidine up to twice the single maximum quantities in the amount required for up to 7 days.

(3) As his own surgery supply the veterinary surgeon shall be allowed only to prescribe on any single day:

1. one of the narcotic drugs specified in paragraph 1 (a) up to the established maximum quantity or one of the narcotic drugs specified in paragraph 1 (b),
2. up to 30 mg alfentanil for pre-medication and anaesthesia including neuroleptoanalgesia,
3. up to 1,000 mg cocaine for surgeries on the eye as
   a) solution up to a content of 20 per cent, or
   b) eye tablet or ointment to a content of 2 per cent,
4. up to 30 mg etorphine for immobilization of animals kept in the zoo , in the circus or in game preserves, to be administered personally or in the presence of the prescribing person,
5. up to 10 mg fentanyl for pre-medication and anaesthesia including neuroleptoanalgesia, for diagnostic surgeries and for immobilization and
6. up to 40,000 mg pentobarbital for pre-medication and anaesthesia as well as for putting down animals.

(4) The veterinary surgeon who heads or supervises a partial unit (ward) of a departmentized veterinary hospital or who heads or supervises an undepartmentized veterinary hospital shall be allowed to prescribe as ward supply for the facility headed or supervised by him,only the narcotic drugs specified in the paragraphs 1 and 3 nos 2,3,5 and 6 in consideration of the restrictions pursuant to paragraph 3 as regards intended purpose, content and dosage form."

4. Section 6 shall be amended as follows:

a) In paragraph 1 no 3 (b) shall be replaced by (b) and (c):

"b) in a finished drug

designation, dosage form, weight quantity of the narcotic drug contained therein per package unit, in separate preparations per dosage unit, and the number of units;

c) in homoeopathic preparations

the degree of dilution of the narocitc drug contained therein and the weight quantity of the package unit, in separate preparations the number the dosage units, in a mixture of several preparations also the weight percentage of the dilution contained in the narcotic drug."

b) In paragraph 1 nos 4 to 9 shall be worded as follows:

"4. the directions for use specifying the single and the daily dose, except in the cases referred to in section 2 paragraphs 2 no 3,

5. in the cases referred to in section 2 paragraph 2 the note saying "quantity veterinarily justified" and, in any case, in addition, the statement as to for how many days the prescription is made,

6. the intended purpose in prescribing alfentanil, cocaine, etorphine and fentanyl as well as buprenorphin and morphine in the cases referred to in section 2

paragraph 2 no 3 and pentobarbital in the cases referred to in section 3 paragraph 2 no 4 and in section 4 paragraph 3 no 6,

7. the name of the prescribing medical practitioner or dental or veterinary surgeon, his professional title and address including phone number, in case of deputizing capacity also the note saying "in deputizing capacity,

8. in the cases referred to in section 2 paragraph 3, section 3 paragraph 2 and section 4 paragraph 3 the note saying "surgery supply" instead of the specifications referred to in nos 1 and 4,

9. in the cases referred to in section 2 paragraph 4, section 3 paragraph 3 and section 4 paragraph 4 the name or the designation and the address of the facility for which the ward supply is intended instead of the specifications referred to in nos 1 and 4,".

c) Paragraph 2 shall be worded as follows:

"(2) The specifications according to paragraph 1 shall be made with an indelible pencil or ball point pen and be identical on all parts of the narcotic drugs prescription form. In this respect, the specifications under the nos 2 to 6, 8 and 10 shall be entered by the prescriber in his own handwriting. Any change of prescription with regard to the specifications under paragraph 1 nos 2 to 6 shall be entered by the prescriber, in his own handwriting, on all parts of the narcotic drugs prescription form and be confirmed by his signature. The specifications under paragraph 1 nos 1, 7 with the exception of the note saying "in deputizing capacity", as well as no 9 may also be entered by another person, in typing or with stamps."

5. Section 8 paragraph 2 sentence 2 shall be worded as follows:

"For merchant ships he may prescribe only the following narcotic drugs:

1. Hydromorphone,
   Opium, standardized,
   Pentazocine and
   Pethidine

   for merchant ships which have no ship's doctor,

2. Hydrocodone,
   Hydromorphone,
   Methamphetamine,
   Morphine,
   Opium, standardized,
   Oxycodone,
   Pentazocine and
   Pethidine

   for merchant ships which have a ship's doctor."

6. Section 10 sentence 1 shall be amended as follows:

   a) In number 2 (a) to (c) shall be worded as follows:

      "2. a) contrary to section 2 paragraph 1 or 2 sentence 1 or section 3 paragraph 1 for a patient,

      b) contrary to section 2 paragraph 3, section 3 paragraph 2 or section 4 paragraph 3 for his surgery supply or

      c) contrary to section 4 paragraph 1 or 2 for an animal".

   b) Number 3 shall be worded as follows:

      "3. contrary to section 2 paragraph 4, section 3 paragraph 3 or section 4 paragraph 4 prescribes

      a) narcotic drugs for institutions other than those specified thereunder,

      b) narcotic drugs other than those specified thereunder or

      c) narcotic drugs specified thereunder disregarding other restrictions."

7. After section 12 the following section 12 a shall be inserted:

"Section 12a

Transitional Provision

The narcotic drugs "thebacon" and "a preparation containing native opium alkaloids in their natural mixing ratio calculated as morphine" may be prescribed by the medical practitioner, the dental or veterinary surgeon until 31 July 1988 pursuant to the provisions valid so far."

Article 3

New Version Authorization

The Federal Minister for Youth, Family Affairs, Women and Health may publish in the Federal Law Gazette the text of Schedules I to III of the Narcotics Act as well as the text of the Narcotic Drugs Prescription Ordinance in the version in force as from 1 August 1986.

Article 4

Berlin Clause

Pursuant to section 14 of the Third Transitory Law in conjunction with section 41 sentence 2 of the Narcotics Act this ordinance shall also apply in the Land of Berlin.

Article 5

Entry into Force

This ordinance shall enter into force on 1 August 1986.

Bonn, 23 July 1986

Dr. Helmut Kohl
Federal Chancellor

Rita Süssmuth
Federal Minister for Youth, Family Affairs, Women and Health